PRELIMINARY AMENDMENT

Serial Number: 09/634207 Filing Date: August 9, 2000

INDOLE COMPOUNDS USEFUL FOR THE TREATMENT OF CANCER

IN THE SPECIFICATION

Please amend the specification as follows:

TECH ON TER INDUSTRIAN SON Please amend the paragraph at page 3, line 15 to page 4, line 4 as follows:

The present invention provides indole compounds of formula (I):

$$(R^6)_n = \begin{cases} R^5 & R^4 \\ R^3 & R^2 \\ R^7 & R^1 & Y-Z \end{cases}$$

wherein R¹ is lower alkyl, lower alkenyl, (hydroxy)lower alkyl, lower alkynyl, phenyl, benzyl or 2-thienyl, R², R³, R⁴ and R⁵ are the same or different and are each hydrogen or lower alkyl; each R⁶ is individually hydrogen, lower alkyl, hydroxy, (hydroxy)lower alkyl, lower alkoxy, benzyloxy, lower alkanoyloxy, nitro or halo, n is 1-3, R⁷ is hydrogen, lower alkyl or lower alkenyl, X is oxy and thio, Y is carbonyl, (CH₂)₁₋₃, (CH₂)₁₋₃C(O), or (CH₂)₁. $_3SO_2$ and Z is $(\omega$ -(4-pyridyl)(C₂-C₄ alkoxy), $(\omega$ -((R⁸)(R⁹) amino)(C₂-C₄ alkoxy), wherein R⁸ and R⁹ are each H, (C₁-C₃)alkyl or together with N are a 5- or 6-membered heterocyclic ring comprising 1-3 N(R⁸), S or nonperoxide O; an amino acid ester of (w-(HO)(C2-C4))alkoxy, N(R8)CH(R8)CO2H, OCH2CH2N(CH3)3+, or 1'-D-glucuronyloxy; or Y-Z is (CH₂)₁₋₃R¹⁰ wherein R¹⁰ is OH, (C₂-C₄)acyloxy, SO₃H, PO₄H₂, N(NO)(OH), SO₂NH₂, PO(OH)NH₂, [[OCH₂CH₂N(CH₂)₂, *₇]] or tetrazolyl; or a pharmaceutically acceptable salt thereof.

Please amend the paragraph at page 8, lines 5 to 23, as follows:

Indole compounds of the present inventions include compounds of formula (I):

EI

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Title: INDOLE COMPOUNDS USEFUL FOR THE TREATMENT OF CANCER

$$(R^{6})_{ri}$$
 R^{5}
 R^{4}
 R^{3}
 R^{2}
 R^{7}
 R^{7}
 R^{7}
 R^{7}
 R^{7}
 R^{7}
 R^{7}
 R^{7}

wherein R¹ is selected from the group consisting of lower alkyl, lower alkenyl, (hydroxy)lower alkyl, lower alkynyl, phenyl, benzyl and 2-thienyl, R², R³, R⁴ and R⁵ are the same or different and are each selected from the group consisting of hydrogen and lower alkyl, each R⁶ is individually selected from the group consisting of hydrogen, lower alkyl, hydroxy, (hydroxy)lower alkyl, lower alkoxy, benzyloxy, lower alkanoyloxy, nitro and halo, n is 1-3, R⁷ is selected from the group consisting of hydrogen, lower alkyl and lower alkenyl, X is selected from the group consisting of oxy and thio, Y is selected from the group consisting of carbonyl (CH₂)₁₋₃, (CH₂)₁. ₃SO₂ or (CH₂)₁₋₃C(O), and Z is selected from the group consisting of hydroxy, lower alkoxy optionally substituted with OH, 4-pyridyl, amino, lower alkylamino, di(lower alkyl)amino, [[er]] OCH2CH2N(CH3)2, N-morpholino; amino, lower alkylamino, [(carboxy)(lower alkyl)]amino, di(lower)alkylamino and phenylamino, or Y-Z is $(CH_2)_{1.3}R^{10}$ wherein R^{10} is OH, (C_2-C_4) acyloxy, SO_3H , PO_4H_2 , N(NO)(OH), SO₂NH₂, PO(OH)NH₂, OCH₂CH₂N(CH₃)₃⁺, or tetrazolyl or a pharmaceutically acceptable salt thereof. Lower alkyl, alkenyl, alkanoyl, etc. indicates a branched, cyclic or straight chain C₁-C₆ group, preferably a C₁-C₄ group, including cycloalkyl and (cycloalkyl)alkyl. (Hydroxy)lower alkyl or alkoxy is preferably 1- or 2hydroxyethyl.

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